UNIVERSITATEA DE MEDICINĂ ȘI FARMACIE
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ANEXA 2

FIȘA DE VERIFICARE A ÎNDEPLINIRII STANDARDELOR MINIMALE ÎN VEDEREA OBȚINERII ATESTATULUI DE ABILITARE ÎN CADRUL I.O.S.U.D. U.M.F. ”GRIGORE T POPA” DIN IAȘI

1. STANDARD MINIMAL ARTICOLE ISI AUTOR PRINCIPAL*

*calitatea de autor principal este atribuită pentru primul autor, ultimul autor, autorul corespondent, sau alți autori a căror contribuție este indicată explicit în cadrul publicației a fi egală cu contribuție primului autor sau a autorului corespondent

Domeniul Medicină, Farmacie: 10 articole, publicate pe parcursul întregii activități

Domeniul Medicină Dentară: 8 articole în reviste cu factor de impact>0.3, de la ultima promovare sau ultimii 5 ani (pentru candidații care provin din afara sistemului de învățământ)

RAPORTARE ARTICOLE ISI AUTOR PRINCIPAL
(listă, cu link-urile corespunzătoare pentru verificare, active)

1. Ciobanu, C. G., Nucă, I., Popescu, R., Antoci, L. M., **Caba, L.,** Ivanov, A. V., Cojocaru, K.-A., Rusu, C., Mihai, C.-T., Pânzaru, M. C. *Narrative Review: Update on the Molecular Diagnosis of Fragile X Syndrome.* Int J Mol Sci 2023, 24(11), 9206. doi: 10.3390/ijms24119206 (contribuții egale). IF=6.208
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2. Daniela Luminita Zob, Iolanda Augustin, **Lavinia Caba***, Monica-Cristina Panzaru, Setalia Popa, Alina Delia Popa, Laura Florea, and Eusebiu Vlad Gorduza. *Genomics and Epigenomics in the Molecular Biology of Melanoma—A Prerequisite for Biomarkers Studies.*Int J Mol Sci 2022, 24 (1): 716. doi: 10.3390/ijms24010716 (autor corespondent). IF=6.208
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4. Laura Florea, **Lavinia Caba*** and Eusebiu Vlad Gorduza. *Genetic Heterogeneity in Bartter Syndrome: Clinical and Practical Importance.* Front Pediatr 2022, June 3;10:908655. doi: 10.3389/fped.2022.908655. eCollection 2022. (autor corespondent). IF=3.569
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5. Monica-Cristina Pânzaru, **Lavinia Caba***, Laura Florea*, Elena Emanuela Braha, Eusebiu Vlad Gorduza. *Epidermolysis Bullosa - a different genetic approach in correlation with genetic heterogeneity.* Diagnostics (Basel) 2022, 12(6), 1325. doi:10.3390/diagnostics12061325 (autor corespondent). IF=3.992
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6. Popescu, R., Grămescu, M., **Caba, L**., Pânzaru, M.C., Butnariu, L., Braha, E., Popa, S., Rusu, C., Cardos, G., Zeleniuc, M., Martiniuc, V., Gug, C., Păduraru, L., Stamatin, M., Diaconu, C.C., Gorduza, E.V. *A Case of Inherited t (4; 10)(q26; q26. 2) Chromosomal Translocation Elucidated by Multiple Chromosomal and Molecular Analyses. Case Report and Review of the Literature.* Genes (Basel) 2021, *12*(12), p.1957, doi: 10.3390/genes12121957. (autor corespondent). IF=4.141
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8. Antohi, C., Haba, D., **Caba, L.**, Ciofu, M.L., Drug, V.L., Bărboi, O.B., Dobrovăț, B.I., Pânzaru, M.C., Gorduza, N.C., Lupu, V.V., Dimofte, D., Gug, C., Gorduza, E.V. *Novel Mutation in APC Gene Associated with Multiple Osteomas in a Family and Review of Genotype-Phenotype Correlations of Extracolonic Manifestations in Gardner Syndrome.* Diagnostics (Basel) 2021, *11*(9), p.1560. doi: 10.3390/diagnostics11091560 (autor corespondent). IF=3.992
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9. Florea, L., **Caba, L.** and Gorduza, E.V. *Bardet–Biedl Syndrome—Multiple Kaleidoscope Images: Insight into Mechanisms of Genotype–Phenotype Correlations.* Genes (Basel) 2021, *12*(9), p.135. doi: 10.3390/genes12091353 (autor corespondent). IF=4.141
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11. **Caba, L**., Gug, C., & Gorduza, E. V. *Heterogeneity in combined immunodeficiencies with associated or syndromic features.* Exp Ther Med 2021, *21*(1), 1-1. doi: 10.3892/etm.2020.9517. IF=2.751.
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13. **Caba L.,** Rusu C., Plăiaşu V., Gug G., Grămescu M., Bujoran C., Ochiană D., Voloşciuc M., Popescu R., Braha E., Pânzaru M., Butnariu L., Sireteanu A., Covic M., Gorduza E.V. *Ring autosomes: some unexpected findings.* Balkan J Med Genet 2012, 15 (2), 35-46. doi: 10.2478/bjmg-2013-0005. IF=0.077.
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15. Pânzaru M., Rusu C., Volosciuc M., Braha E., Butnariu L., Gramescu M., Popescu R., **Caba L.,** Bujoran C., Ivanov I., Macovei M., Sireteanu A., Covic M., Gorduza E.V. *Benefits of cytogenetic testing in diagnosis of plurimalformative syndromes with congenital heart defects.* Rev Rom Med Lab 2012, 20 (3/4) 265-272 (autor corespondent). IF=0.097
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2. STANDARD MINIMAL ARTICOLE ISI COAUTOR
Domeniul Medicină, Farmacie: 5 articole, publicate pe parcursul întregii activităţi

RAPORTARE ARTICOLE ISI COAUTOR
(listă, cu link-urile corespunzătoare pentru verificare, active)

1. Khan, S., Focșa, I. O., Budișteanu, M., Stoica, C., Nedelea, F., Bohîlțea, L., **Caba, L.**, Butnariu, L., Panzaru, M., Rusu, C., Jurca, C., Chirita-Emandi, A., Banescu, C., Abbas, W., Sadeghpour, A., Baig, S. M., Balgradean, M., Davis, E. E. Exome sequencing in a Romanian Bardet-Biedl syndrome cohort revealed an overabundance of causal BBS12 variants. *Am J Med Genet 2023,* 1–16. doi: 10.1002/ajmg.a.63322. IF=2.578

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2. Popa, A. D., Niță, O., Gherasim, A., Enache, A. I., **Caba, L.**, Mihalache, L., & Arhire, L. I. A Scoping Review of the Relationship between Intermittent Fasting and the Human Gut Microbiota: Current Knowledge and Future Directions. *Nutrients* 2023, *15*(9), 2095. doi: 10.3390/nu15092095. IF=6.706

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3. Panzaru, M. C., Florea, A., **Caba, L.**, & Gorduza, E. V. Classification of osteogenesis imperfecta: Importance for prophylaxis and genetic counseling. *World J Clin Cases* 2023, *11*(12), 2604-2620. doi: 10.12998/wjcc.v11.i12.2604. IF=1.534
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4. Ungureanu, M.-C.; Hrisca, A.; **Caba, L.**; Teodoriu, L.; Bilha, S.; Preda, C.; Leustean, L. *SHOX Deletion and Idiopathic Short Stature: What Does the Clinician Need to Know? Case Series Report.* Diagnostics (Basel) 2023, *13*, 105. doi:10.3390/diagnostics13010105. IF=3.992

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5. Gug, C., Mozos, I., Ratiu, A., Tudor, A., Gorduza, E.V., **Caba, L.**, Gug, M., Cojocariu, C., Furau, C., Furau, G., Vaida, M.A., Stoicanescu, D. *Genetic Counseling and Management: The First Study to Report NIPT Findings in a Romanian Population.* Medicina (Kaunas) 2022, *58*, 79. doi:10.3390/medicina58010079. IF=2.948

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6. Resmerita, I., Cozma, R.S., Popescu, R., Radulescu, L.M., Panzaru, M.C., Butnariu, L.I., **Caba, L.,** Ilie, O.D., Gavril, E.C., Gorduza, E.V. and Rusu, C. *Genetics of Hearing Impairment in North-Eastern Romania—A Cost-Effective Improved Diagnosis and Literature Review.* Genes (Basel) 2020, *11*(12), p.1506, doi: 10.3390/genes11121506. IF=4.096.

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7. Nemescu, D., Constantinescu, D., Gorduza, V., Carauleanu, A., **Caba, L.** and Navolan,

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8. Sireteanu, A., Popescu, R., Braha, E.E., Bujoran, C., Butnariu, L., **Caba, L**., Graur, E., Gorduza, E.V., Grămescu, M., Ivanov, I.C. and Pânzaru, M. *Detection of chromosomal imbalances using combined MLPA kits in patients with syndromic intellectual disability.* Rev Rom Med Lab 2014, *22*(2). pp.157-164.doi:10.2478/rrlm-2014-0019.IF=0.239. https://www.rrml.ro/articole/2014/2014_2_1.pdf
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9. Gorduza E.V**.**, Popescu R., **Caba L**., Ivanov I., Martiniuc V., Nedelea F., Militaru M., Socolov D.G. *Prenatal diagnosis of 21 trisomy by quantification of methylated foetal DNA in maternal blood: study on 10 pregnancies (Diagnosticul prenatal al trisomiei 21 prin cuantificarea ADN-ului fetal metilat din sângele matern: studiu pe 10 sarcini).* Rev Rom Med Lab 2013, 21 (3/4) 275-284. doi: 10.2478/rrlm-2013-0030. IF=0.171.
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3. STANDARD MINIMAL FACTOR CUMULAT DE IMPACT*

*se calculează ca sumă a factorilor de impact a revistelor în care au fost publicate articolele **în calitate de autor principal**, **în anul aparţiei articolului**
Domeniul Medicină, Farmacie: 10, pe parcursul întregii activităţi
Domeniul Medicină Dentară: 2,4, de la ultima promovare sau ultimii 5 ani (pentru candidaţii care provin din afara sistemului de învăţământ)

RAPORTARE FACTOR CUMULAT DE IMPACT

51.795

(cifră, calculată conform indicaţiilor, cu link-urile corespunzătoare pentru verificare, active)
Factor cumulat de impact= 6.208+ 6.208+ 2.919+ 3.569+ 3.992+ 4.141+ 6.064+ 3.992+ 4.141+ 2.948+ 2.751+ 3.688+ 0.077+ 1+ 0.097=51.795

4. STANDARD MINIMAL INDICE HIRSCH*

Domeniul Medicină, Medicină Dentară, Farmacie = 6
*conform ISI Web of Science, Core Collection, Thomson Reuters

RAPORTARE INDICE HIRSCH
(cifră, cu link-urile corespunzătoare pentru verificare, active)

Indice HIRSCH **7**

Indice HIRSCH pdf
https://www.webofscience.com/wos/woscc/citation-report/8e8a8751-fc2f-4fed-8d9f-73e975d8a4c0-85677bb1?state=%7B%7D

5. STANDARD MINIMAL ARTICOLE BDI*

Domeniul Medicină Dentară: 20 articole de la ultima promovare, în calitate de autor principal *echivalare = 1 articol ISI autor principal =3 articole BDI autor principal, dar nu invers
Nu este cazul

25.06.2023 Conf. univ. dr. Lavinia Caba